Listing of Claims:

- 1. (Cancelled) A pharmaceutical composition for therapeutic or prophylactic use comprising a silica containing solid having an average particle size of about 6 microns or less.
- 2. (Cancelled) The pharmaceutical composition according to claim 1 wherein the silica containing solid is selected from the group consisting of zeolites, silicas, clays, double hydroxides, and mixtures thereof.
- 3. (Cancelled) The pharmaceutical composition according to claim 1 wherein the silica containing solid is zeolite containing encapsulated metals or metal complexes.
- 4. (Cancelled) The pharmaceutical composition according to claim 3 wherein the metal complexes are metal salen complexes, phthalocyanines, corrinoides or porphyrines.
- 5. (Cancelled) The pharmaceutical composition according to claim 1 wherein the silica containing solid is silica gel or other silicas containing encapsulated metals, metal complexes, proteins, DNA or whole cells or tissue samples.
- 6. (Cancelled) The pharmaceutical composition according to claim 1 wherein the silica containing solid is mesoporous aluminosilicate containing encapsulated metal complexes, proteins, DNA or small molecules having pharmaceutical activity.
- 7. (Cancelled) The pharmaceutical composition according to claim 1 wherein the silica containing solid is modified by surface adsorption of molecules to enhance the bioavailability of the silica containing solid.
- 8. (Cancelled) The pharmaceutical composition according to claim 7 where the silica containing solid is modified by surface adsorption of molecules selected from the group consisting of vitamin B12 and silanes.

- 9. (Cancelled) The pharmaceutical composition according to claim 1 where the silica-containing solid is dealuminated.
- 10. (Cancelled) The pharmaceutical composition according to claim 1 where the pores of the silica containing solid are modified by silanation, methylation, surfactant adsorption or other chemical reaction to change the wettability, charge or size of the pores.
- 11. (Cancelled) A method to modify gene expression, cell proliferation, death, growth rate or differentiation by administering to a mammal a silica containing solid as an antioxidant or oxidant.
- 12. (Cancelled) A method to enhance immunogeneity of protein antigens, other biological macromolecules, whole cells or cell fragments by administering to a mammal in need thereof a silica containing solid as a vaccine adjuvant in combination with protein antigens, whole cells or cell fragments.
- 13. (Cancelled) A method for providing sustained delivery of a pharmaceutically active agent by using a silica containing solid as a reservoir for the pharmaceutically active agent.
- 14. (Cancelled) The method of claim 13 wherein the pharmaceutically active agent is selected from the group consisting of metals, metal complexes, small molecules, proteins, DNA, cell fragments and whole cells.
 - 15. (New) A method of treating cancer in a patient comprising:administering to the patient a pharmaceutical composition comprising a zeolite.
- 16. (New) The method of claim 1 wherein the zeolite has an average particle size of about 6 microns or less.
 - 17. (New) The method of claim 1 wherein the zeolite is clinoptilolite.
 - 18. (New) The method of claim 3 wherein the clinoptilolite has a mean particle size of 250 nm.
- 19. (New) The method of claim 1 wherein the pharmaceutical composition further comprises at least one of pro-oxidant metal complexes, zinc, silver, cytokines, cells, and tumor antigens.

- 20. (New) The method of claim 5 wherein the cells are live vaccine cells.
- 21. (New) The method of claim 5 wherein the cytokines is IL-12, GM-CSF or interferon gamma.
- 22. (New) The method of claim 1 wherein the cancer is selected from the group consisting of lung cancer and colorectal cancer.
- 23. (New) The method of claim 1 wherein the pharmaceutical composition is administered by injection.